DEPARTMENT OF HEALTH AND HUMAN SERVICES,

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Food and Drug Administration

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[Docket No. 02N-0109]

Agency Information Collection Activities; Announcement of OMB Approval; Dissemination of Information on Unapproved/New Uses for Marketed Drugs, Biologics, and Devices

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Dissemination of Information on Unapproved/New Uses for Marketed Drugs, Biologics, and Devices" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (the PRA).

FOR FURTHER INFORMATION CONTACT: Karen L. Nelson, Office of Information Resources Management (HFA-250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-1482.

SUPPLEMENTARY INFORMATION: In the Federal Register of September 16, 2002 (67 FR 58429), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under section 3507 of the PRA (44 U.S.C. 3507). An agency may not conduct or sponsor and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0390. The approval expires on November 30, 2005. A copy of the supporting statement oc02328

for this information collection is available on the Internet at http://www.fda.gov/ohrms/dockets.

Dated: 12 26 02 December 26, 2002.

Margaret M. Dotzel,

Assistant Commissioner for Policy.

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